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USPT	11 and 13	2	<u>L5</u>	
USPT	11 and 12	0	<u>L4</u>	
USPT	melanoma	7496	<u>L3</u>	
USPT lung near5 (cancer or carcinoma)		6496	<u>L2</u>	
USPT	chlorotoxin	7	<u>L1</u>	

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☐ 1. Document ID: US 6028174 A

L5: Entry 1 of 2

File: USPT

Feb 22, 2000

US-PAT-NO: 6028174

DOCUMENT-IDENTIFIER: US 6028174 A

TITLE: Method of diagnosing and treating gliomas

Full Title Citation Front Review Classification Date Reference Claims KMC Draw Desc Image

☐ 2. Document ID: US 5905027 A

L5: Entry 2 of 2

File: USPT

May 18, 1999

US-PAT-NO: 5905027

DOCUMENT-IDENTIFIER: US 5905027 A

TITLE: Method of diagnosing and treating gliomas

Full Title Citation Front Review Classification Date Reference Claims KMC Draw Desc Image

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Terms	Documents
l1 and l3	2

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20 Documents, starting with Document: 2

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L1
            74 S CHLOROTOXIN
L2
          38153 S SMALL(W) CELL(6A) CARCINOMA
L3
             1 S L1 AND L2
T.4
        148410 S MELANOMA
L5
             8 S L1 AND L4
L6
             3 DUP REM L5 (5 DUPLICATES REMOVED)
=> d bib ab 13
    ANSWER 1 OF 1 CAPLUS COPYRIGHT 2001 ACS
L3
    2000:756551 CAPLUS
ΑN
    133:307331
DN
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    Diagnosis and treatment of neuroectodermal tumors
    Sontheimer, Harald J.; Lyons, Susan A.
IN
PΑ
    UAB Research Foundation, USA
SO
    PCT Int. Appl., 56 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
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    PATENT NO.
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PRAI US 1999-296031
                     19990421
    The present invention provides fusion proteins for the detection and
    treatment of neuroectodermal tumors. Previous work demonstrated that
    chlorotoxin is specific for glial-derived or meningioma-derived
    tumor cells. The current invention has extended the use of
    chlorotoxin-cytotoxin fusion proteins to treat the whole class
    neuroectodermal tumors such as gliomas, meningiomas, ependymonas,
    medulloblastomas, neuroblastomas, gangliomas, pheochromocytomas,
    melanomas, PPNET's, small cell carcinoma of
    the lung, Ewing's sarcoma, and metastatic tumors in the brain. Also,
    diagnostic methods are provided for screening neoplastic neuroectodermal
    tumors.
RE.CNT
       3
RE
(1) Soroceanu; J Neuroscience 1999, V19(14), P5942 CAPLUS
(2) Ullrich; Am J Physiol 1996, V270, PC1511 CAPLUS
(3) Ullrich; Neuroscience 1998, V83(4), P1161 CAPLUS
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ANSWER 1 OF 3 CAPLUS COPYRIGHT 2001 ACS
L6
     2000:756551 CAPLUS
ΑN
DN
     133:307331
ΤI
     Diagnosis and treatment of neuroectodermal tumors
IN
     Sontheimer, Harald J.; Lyons, Susan A.
PA
     UAB Research Foundation, USA
     PCT Int. Appl., 56 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                            DATE
     PATENT NO.
                      KIND
                                           APPLICATION NO.
                                                             DATE
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PΙ
     WO 2000062807
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             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRAI US 1999-296031
                      19990421
     The present invention provides fusion proteins for the detection and
     treatment of neuroectodermal tumors. Previous work demonstrated that
     chlorotoxin is specific for glial-derived or meningioma-derived
     tumor cells. The current invention has extended the use of
     chlorotoxin-cytotoxin fusion proteins to treat the whole class
     neuroectodermal tumors such as gliomas, meningiomas, ependymonas,
     medulloblastomas, neuroblastomas, gangliomas, pheochromocytomas,
     melanomas, PPNET's, small cell carcinoma of the lung, Ewing's
     sarcoma, and metastatic tumors in the brain. Also, diagnostic methods
are
     provided for screening neoplastic neuroectodermal tumors.
RE.CNT 3
(1) Soroceanu; J Neuroscience 1999, V19(14), P5942 CAPLUS
(2) Ullrich; Am J Physiol 1996, V270, PC1511 CAPLUS
(3) Ullrich; Neuroscience 1998, V83(4), P1161 CAPLUS
     ANSWER 2 OF 3 MEDLINE
                                                         DUPLICATE 1
L6
AN
     96396940
                  MEDLINE
DN
     96396940
     Human astrocytoma cells express a unique chloride current.
ΤI
     Ullrich N; Gillespie G Y; Sontheimer H
ΑU
     Neurobiology Research Center, University of Alabama at Birmingham 35294,
CS
     USA.
NC
     RO-1 NS31234 (NINDS)
     P50 HD32901 (NICHD)
     P20 NS31096 (NINDS)
     NEUROREPORT, (1996 Apr 10) 7 (5) 1020-4.
SO
     Journal code: A6M. ISSN: 0959-4965.
CY
     ENGLAND: United Kingdom
     Journal; Article; (JOURNAL ARTICLE)
DT
LA
     English
     Priority Journals
FS
     199702
EM
EW
     19970204
     Human astrocytoma cells were studied using whole-cell patch-clamp
AB
     recording. Voltage-dependent outwardly-rectifying anion currents were
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identified in primary cultures of six freshly resected human brain tumors and in seven established anaplastic astrocytoma/glishlastoma cell lines (U251MG, CH235MG, B73MG, U105MG, D54MG, SK-MG-1, 1 STTG1). Anion currents were not observed in normal, non-neoplastic glial cells, nor in human tumor-derived cells of non-glial origin (melanoma, breast cancer, neuroblastoma, rhabdomyosarcoma). Currents activated at potentials

> 50 mV and showed large transients upon termination of voltage steps. Currents reversed at the predicted equilibrium potential for chloride ions

and could also be recorded when Cl- was replaced by F-, Br- or I-. Currents were inhibited by the Cl- channel blockers **chlorotoxin**, DIDS, and DNDS. These Cl- currents may play a role in the growth control of astrocytoma cells.

L6 ANSWER 3 OF 3 MEDLINE

DUPLICATE 2

AN 96352227 MEDLINE

DN 96352227

TI Human astrocytoma cells express a unique chloride current.

AU Ullrich N; Gillespie G Y; Sontheimer H

CS Interdepartmental Neuroscience Program, Yale University School of Medicine, New Haven, CT 06510, USA.

NC RO-1 NS31234 (NINDS) P50 HD32901 (NICHD) P20 NS31096 (NINDS)

SO NEUROREPORT, (1995 Dec 29) 7 (1) 343-7. Journal code: A6M. ISSN: 0959-4965.

CY ENGLAND: United Kingdom

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 199612

AB Human astrocytoma cells were studied using whole-cell patch-clamp recording. Voltage-dependent outwardly-rectifying anion currents were identified in primary cultures of six freshly resected human brain tumors and in seven established anaplastic astrocytoma/glioblastoma cell lines (U251MG, CH235MG, U373MG, U105MG, D54MG, SK-MG-1, and STTG1). Anion currents were not observed in normal, non-neoplastic glial cells, nor in human tumor-derived cells of non-glial origin (melanoma, breast cancer, neuroblastoma, rhabdomyosarcoma). Currents activated at

> 50 mV and showed large transients upon termination of voltage steps. Currents reversed at the predicted equilibrium potential for chloride

and could also be recorded when Cl- was replaced by F-, Br- or I-. Currents were inhibited by the Cl- channel blockers **chlorotoxin**, DIDS, and DNDS. These Cl- currents may play a role in the growth control of astrocytoma cells.